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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

Applicant: CRAWLEY et al.

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Page 1 of 1

Examiner: Unassigned

Group Art Unit: 1614

U.S. PATENT DOCUMENTS

Examiner's Initials*	Document Number	Date MM/YYYY	Name (Family Name of First Inventor)	Class	Sub Class	Filing Date (if appropriate)
	AR					
	BR					
	CR					
	DR					
	ER					
	FR					

FOREIGN PATENT DOCUMENTS

		Document Number	Date MM/YYYY	Country	Inventor Name	English Abstract		Translation Readily Available	
						Enclosed	No	Enclose	No
TNT	GR	WO 97/03069	01/1997	WIPO	Cockerill et al.				
	HR	WO 98/38984	09/1998	WIPO	Shenoy et al.				
	IR	WO 98/50047	11/1998	WIPO	Liang et al.				
	JR	WO 98/50370	11/1998	WIPO	Tang et al.				
TNT	KR	WO 99/09024	02/1999	WIPO	Chan et al.				
	LR								
	MR								
	NR								

OTHER (Including in this order Author, Title, Periodical Name, Date, Pertinent Pages, etc.)

TNT	OR	Gibson et al.; "Epidermal Growth Factor Receptor Tyrosine Kinase: Structure-Activity Relationships and Antitumour Activity of Novel Quinazolines"; Bioorganic & Medicinal Chemistry Letters, 1997, Vol. 7, No. 21, pp. 2723-2728	TECH CENTER 1600/2900	JUN 14 2002	RECEIVED
	PR	Hong et al.; "Synthesis and Biological Activities of Some N ⁴ -Substituted 4-Aminopyrazolo[3,4-d-pyrimidines"; Journal of Medicinal Chemistry, 1976, Vol. 19, No. 4, pp. 555-558			
	QR	Myers et al.; "The Preparation and SAR of 4-(Anilino), 4-(Phenoxy), and 4-(Thiophenoxy)-Quinazolines: Inhibitors of p56 ^{lck} and EGF-R Tyrosine Kinase Activity"; Bioorganic & Medicinal Chemistry Letters, 1997, Vol. 7, No. 4, pp. 417-420			
TNT	RR	van Muijlwijk-Koezen et al.; "Isoquinoline and Quinazoline Urea Analogues as Antagonists for the Human Adenosine A ₃ Receptor"; J. Med. Chem., 2000, Vol. 43, pp. 2227-2238			
	SR				
	TR				

Examiner

[Signature]

Date Considered:

7/28/03

*EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP § 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to Applicant.